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(56) **References Cited**

U.S. PATENT DOCUMENTS

4,377,446 A	3/1983	Albery
4,525,266 A	6/1985	Schmidt et al.
4,554,383 A	11/1985	Knifton

(Continued)

FOREIGN PATENT DOCUMENTS

DE	196 22 930 A1	12/1997
DE	102 45 337 A1	4/2004

(Continued)

OTHER PUBLICATIONS

Alnashif, Inas M., et al., Electrochemical Generation of Superoxide in Room-Temperature Ionic Liquids, *Electrochem. Solid-State Lett.*, Nov. 2001, pp. D16-D18, vol. 4, No. 11.

(Continued)

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H01M 2300/0022 (2013.01); *H01M 2300/0025*
(2013.01); *Y02E 60/50* (2013.01)

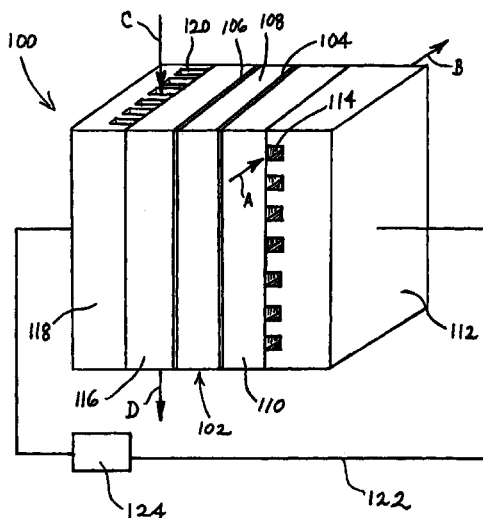
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(57) **ABSTRACT**

An electrochemical cell for applications such as electrochemical fuel cells, or electrochemical cell gas sensors used for detection of target gas species in environments containing or susceptible to presence of same. The electrochemical cell utilizes an ionic liquid as an electrolyte medium, thereby achieving a broader range of operational temperatures and conditions, relative to electrochemical cells utilizing propylene carbonate or other conventional electrolytic media.

15 Claims, 2 Drawing Sheets



(56)

References Cited

U.S. PATENT DOCUMENTS

4,587,003	A	5/1986	Tantram et al.	
5,171,649	A	12/1992	Vaughn	
5,208,112	A *	5/1993	Ludwig et al.	429/419
5,338,429	A *	8/1994	Jolson et al.	204/415
5,683,832	A *	11/1997	Bonhote et al.	429/111
5,846,670	A *	12/1998	Watanabe et al.	429/530
5,855,809	A *	1/1999	Angell et al.	252/62.2
5,865,973	A	2/1999	Kiesele et al.	
5,965,054	A *	10/1999	McEwen et al.	252/62.2
6,667,128	B2	12/2003	Edlund	
7,060,169	B2	6/2006	Röhl	

FOREIGN PATENT DOCUMENTS

EP	0 531 745	A2	3/1993	
EP	0531745		3/1993 G01N 27/416
WO	WO-00/15594	A1	3/2000	
WO	WO-00/16902	A1	3/2000	

OTHER PUBLICATIONS

Cai, Qi, et al., Studies on a sulfur dioxide electrochemical sensor with ionic liquid as electrolyte, Huadong Shifan Xuebao, Ziran Kexueban, 2001, pp. 57-60, vol. 3.

Carter, Michael T., et al., Electrochemical reduction of dioxygen in room-temperature imidazolium chloride-aluminum chloride molten salts, Inorg. Chem., Mar. 6, 1991, pp. 1149-1151, vol. 30, No. 5.

Seddon, K.R., Room-Temperature Ionic Liquids: Neoteric Solvents for Clean Catalysis, 1996, pp. 693-697, vol. 37, No. 5.

Seddon, K.R., Ionic liquids for clean technology: An update, Molten Salt Forum, 1998, pp. 53-62, vol. 5-6.

Wang, Rong, et al., O₂ Gas Sensor Using Supported Hydrophobic Room-temperature Ionic Liquid Membrane-coated electrode, Chem. Lett., Jan. 2004, pp. 6-7, vol. 33, No. 1.

Wang, Rong, et al. A Novel Amperometric O₂ Gas Sensor Based on Supported Room-Temperature Ionic Liquid Porous Polyethylene Membrane-Coated, Electroanalysis, Jan. 2004, pp. 66-72, vol. 16, No. 1-2.

Cai Qi, Xian Yue-zhong, Li Hui, Zhang Yang-ming, Tang Jie, Jin Li-tong (Center of Chemistry of Ionic Liquids, East China Normal University, Shanghai 200062, China), Study on Sulfur Dioxide Electrochemical Sensor Using an Ionic Liquid, Journal of East China Normal University (Natural Science) No. 3, Sep. 2001, pp. 1-7, Article No. 1000-5641 (2001) 03-0057-04. (English Abstract).

Extended European Search Report corresponding to European Application No. 11 15 5949 dated Aug. 30, 2011.

English translation of abstract of EP 0531745 (A2).

English translation of abstract of DE 19622930 (A1).

English translation of abstract of DE 10245337 (A1).

European Patent Office's Jul. 23, 2013 Office Action corresponding to Application No. 11 155 949.8.

Knake R et al., "Amperometric sensing in the gas-phase", Analytica Chimica Acta, Elsevier, Amsterdam, NL, vol. 549, No. 1-2, Sep. 6, 2005, pp. 1-9.

Advances in Electrochemistry and Electrochemical Engineering, vol. 10, John Wiley & Sons, 1976.

* cited by examiner

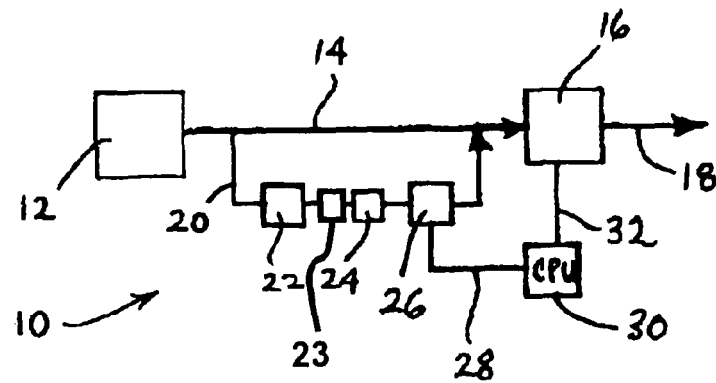


FIG. 1

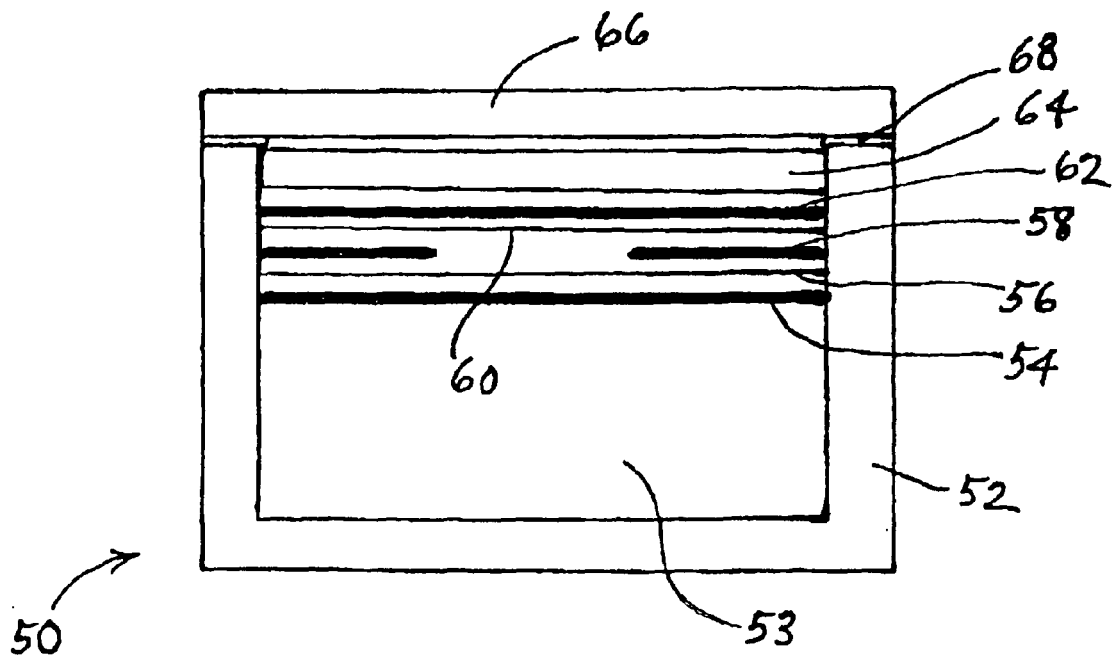


FIG. 2

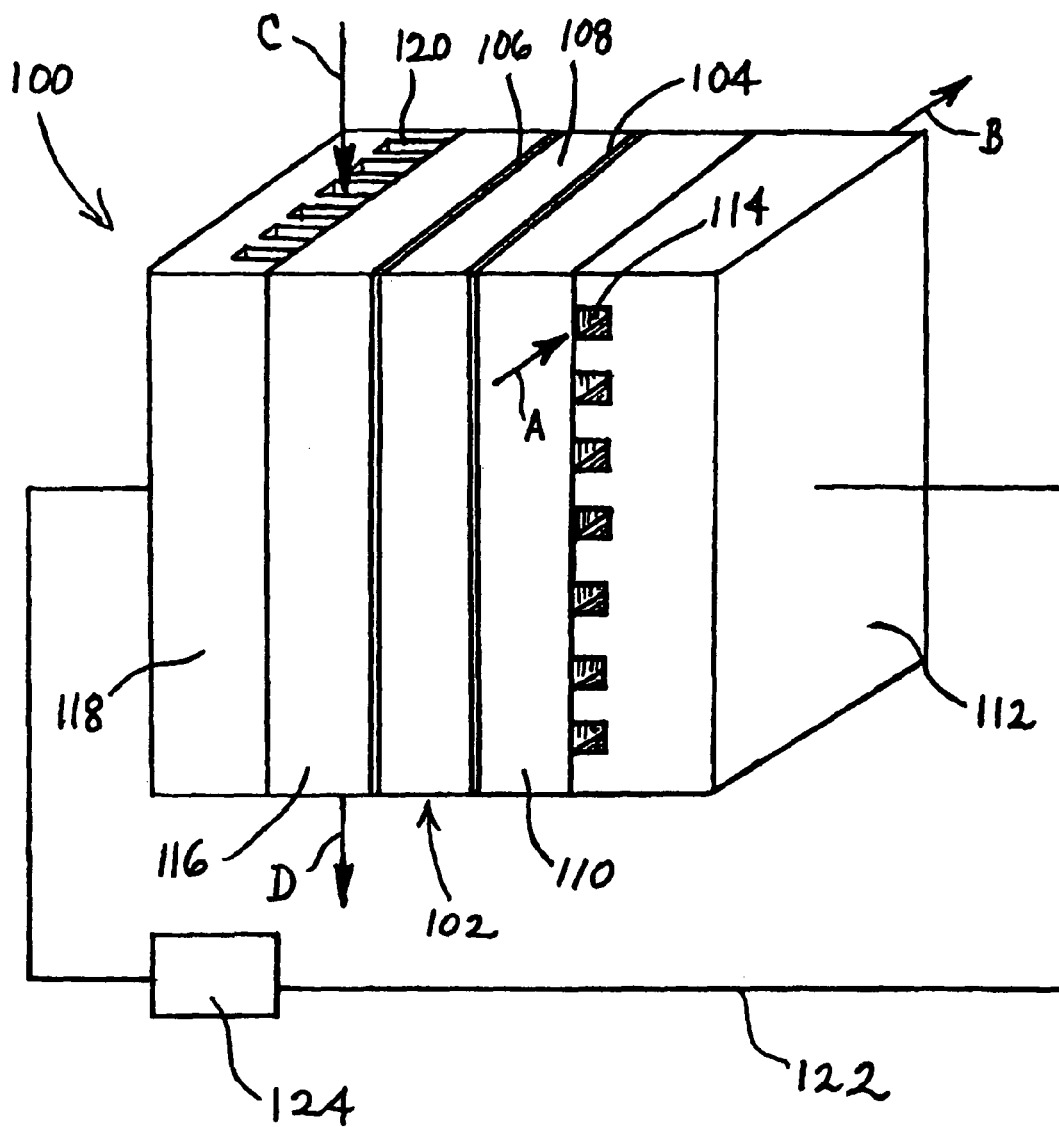


FIG. 3

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ELECTROCHEMICAL CELLS, AND GAS SENSOR AND FUEL CELL DEVICES COMPRISING SAME

CROSS-REFERENCE TO RELATED APPLICATIONS

This is a divisional of U.S. patent application Ser. No. 10/218,262 filed Aug. 14, 2002 in the name of Andreas Röhl for "Electrochemical cells, and gas sensor and fuel cell devices comprising same," which will issue on Jun. 13, 2006 as U.S. Pat. No. 7,060,169, the entirety of which is incorporated herein by reference for all purposes. Priority of said U.S. patent application Ser. No. 10/218,262 is claimed under the provisions of 35 USC 120.

BACKGROUND OF THE INVENTION

1. Field of the Invention

The present invention relates generally to electrochemical cells and to devices incorporating same, such as fuel cells, sensors, and electrochemical cell gas sensors used for detection of gases in environments containing or susceptible to presence of same, and in a specific aspect, the invention relates to an electrochemical cell utilizing ionic liquid as an electrolyte medium thereof.

2. Description of the Related Art

Gas sensors are used in many commercial and industrial applications, including workplace monitoring for the presence of toxic or otherwise hazardous or deleterious gases and in other applications where health and safety issues require detection of specific gases in the ambient environment.

In these various applications, it is frequently necessary or desirable to monitor concentration of selected gas species down to levels on the order of a few parts per million and less.

Gas sensors used in the foregoing applications include electrochemical gas sensors, which may operate to electrochemically reduce the gas species to be monitored. Alternatively, the gas sensor may operate by electrochemically oxidizing the target gas species sought to be detected. As a still further alternative, the electrochemical gas sensor may operate by indirect oxidation or reduction reaction of a compound that is produced in the gas sensor device, involving the target gas to be detected in the monitored gaseous environment.

Electrochemical gas sensors utilize sensor cells that typically contain three electrodes—the working electrode, the reference electrode and the counter electrode, although gas sensor cells are known having 2-electrode and 4-electrode structures. The electrodes are conventionally mounted within a housing that additionally contains an electrolyte, contacts and electrical wires forming electronic circuitry of the sensor, and a gas permeable membrane that keeps the electrolyte within the cell and allows the gas to contact the measuring electrode.

Electrochemical sensor cells require an electrolyte as a component of the electrochemical cell. The electrolyte performs the transport of electrical charge between the different electrodes and therefore enables an electrical current to flow. The transport of electrical charge by the electrolyte is ionic in character rather than involving charge transport by electrons.

The electrolyte usually is constituted by a liquid solvent containing electrolyte/salt component(s), wherein the solvent comprises water, or alternatively a non-aqueous solvent medium, but the electrolyte may otherwise comprise a solid electrolyte such as yttrium-stabilized zirconia (YSZ) for usage at high temperatures of about 450 to 950° C. or an ion exchange membrane such as a Nafion® membrane (commer-

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cially available from DuPont de Nemours and Company, Wilmington, Del.) that is saturated with water.

Liquid electrolytes in some instances can be 'solidified' by addition thereto of gel-forming agents, but the resultantly solidified electrolyte materials still rely on the presence of liquid solvent medium for operability, as do ion exchange systems such as the ion exchange membranes mentioned above.

Gas sensors are operationally "open systems" in the sense of requiring gas flow therethrough for detection of gas component(s) in the gas stream with which the gas sensor is contacted. One major deficiency of conventional gas sensors is the limited service life of liquid electrolytes that include gels or wetted membranes, since these materials dry out over time as solvent evaporates therefrom.

Water-based gas sensor systems are designed to be in equilibrium with ambient humidity of the environment being monitored, but their utility is generally restricted to a specific humidity range.

Solid electrolyte gas sensors are difficult to construct for efficient operation, since diffusion of gas in the solid electrolyte material is hindered in relation to liquid electrolytes, there is no transport of reactants in the solid electrolyte, and known solid electrolyte materials tend to be hygroscopic and thus are difficult to stabilize in humid environments. Additionally, solid electrolytes (i.e. solid materials providing intrinsic ion transport by usually oxide ions O^{2-}) need high operating temperatures. For example, yttrium-stabilized zirconia (YSZ) used in oxygen sensors does not provide sufficient ion conductivity at temperatures below 450° C.

The art therefore continues to seek improvements in electrochemical cell gas sensors.

SUMMARY OF THE INVENTION

The present invention relates generally to electrochemical cells, for applications such as electrochemical energy supplies (fuel cells, batteries, etc.) and electrochemical cell gas sensors.

In one aspect, the invention relates to an electrochemical cell including an electrolyte therein in electrical contact with electrochemical cell electrodes, wherein the electrolyte comprises an ionic liquid.

In another aspect, the invention relates to a method of increasing performance of an electrochemical cell, comprising using in such cell an electrolytic medium including an ionic liquid electrolyte.

Other aspects, features and advantages of the invention will be more fully apparent from the ensuing disclosure and appended claims.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 is a schematic representation of a process system employing an electrochemical cell gas sensor according to the present invention, in an illustrative embodiment thereof.

FIG. 2 is a schematic representation of an electrochemical cell gas sensor and filter unit according to one embodiment of the invention.

FIG. 3 is a simplified schematic representation of an electrochemical fuel cell according to another embodiment of the invention.

DETAILED DESCRIPTION OF THE INVENTION, AND PREFERRED EMBODIMENTS THEREOF

The present invention relates to electrochemical cells utilizing ionic liquids as electrolyte media.

As used herein, the term "ionic liquid" means a salt comprising at least one cation and one anion and being liquid at temperatures below 250° C., preferably below 100° C., and most preferably below 50° C. (e.g., 20-30° C.).

The ionic liquids employed in the broad practice of the invention may be of any suitable type, and include those disclosed in: International Publication No. WO 00/16902 for "IONIC LIQUIDS;" International Publication No. WO 00/15594 for "PROCESS FOR MAKING AROMATIC ALDEHYDES USING IONIC LIQUIDS;" U.S. Pat. No. 4,554,383; Seddon, K. R., Molten Salt Forum, 5-6, pp. 53-62 (1998); and Seddon, K. R., Kinetics and Catalysts, 37, 5, pp. 743-748 (1996), the disclosures of which hereby are incorporated herein by reference in their respective entireties.

The electrochemical cell devices of the present invention include electrochemical cell gas sensors and fuel cells, wherein ionic liquid is employed as an electrolyte medium, having utility in many environments including but not limited to:

- a) environments having broad humidity range, preferably environments having low relative humidity;
- b) environments having high air flow; and
- c) environments operating at higher temperatures.

Additionally, the electrochemical cell devices of the present invention include electrochemical cell gas sensors, having potentially smaller physical dimensions as compared to current electrochemical cell gas sensing devices using aqueous and organic solvent electrolyte. Such smaller dimensions are possible, because unlike current electrochemical sensors, solvent evaporation issues are substantially non-existent allowing for reduced amounts of electrolyte and elimination of reservoir(s) for solvent replacement.

By minimizing the evaporation issue, the electrochemical cell gas sensors of the instant invention are advantageously available for detecting a broader range of target compounds and sensor lifetime is no longer restricted by the amount of electrolyte.

In one aspect, the electrochemical cell of the invention comprises an electrochemical cell gas sensor, in which an ionic liquid is employed as an electrolyte. The electrochemical cell gas sensor may be of any suitable fabrication or construction, e.g., comprising a housing, with two or more electrodes operatively arranged in an interior volume of the housing and communicating with electrical circuitry means (e.g., contacts, wires, conductive films, circuit components, etc.) and the ionic liquid electrolyte.

The ionic liquid may be of any suitable type and may have a melting point that is \leq 250° C. The melting point of the ionic liquid may be above or below the operating temperature of a particular process.

The vapour pressure of the ionic liquids of the instant invention at room temperature is negligible to non-measurable, as compared to electrolyte materials of the prior art (such as water, acrylonitrile, propylene carbonate, and the like).

Illustrative ionic liquids useful as electrolyte materials in electrochemical cell gas sensors in the practice of the present invention include, without limitation, salts with imidazolium or pyridinium cations. The cations that may be usefully employed in a given application of the invention include cations with alkyl and/or aryl substituents, optionally wherein the alkyl and/or aryl substituents are themselves further substituted, e.g., with halo, loweralkyl (C_1 - C_4), hydroxyl, amine, etc.), such cations being capable of forming salts when reacted with corresponding anions.

Illustrative anions for such purpose include, without limitation, halides (chlorides, iodides, bromides, fluorides),

nitrate, nitrite, tetrafluoroborate, hexafluorophosphate, trifluoromethanesulfonates and other polyfluoroalkanesulfonates, e.g., nonaflate, bis(trifluoromethylsulfonyl)imides, methylsulfates, acetates, fluoroacetates and other anions of fluoroalkanoic acid.

In one embodiment, the ionic liquid electrolyte comprises at least one salt selected from the group consisting of salts with the general formula C^+A^- where C^+ represents a quaternary ammonium and/or phosphonium salts and A^- represents any known anion that can form ionic liquids in the meaning described in the present invention. Non-limiting examples include methyl-octyl-imidazolium-chloride and butyl-methyl-imidazolium-bis-trifluoromethane-sulfonimide.

The low vapour pressure ionic liquids in gas sensor electrochemical cells of the present invention may include compounds as described above, in single salt electrolyte compositions, as well as use of same and similar compounds in mixtures together with varying amounts of aluminum halides, preferably aluminum chloride, or other salts that increase the conductivity of the electrolytic medium, or otherwise enhance the gas detection sensitivity of the gas sensor electrochemical cell.

The electrochemical cell gas sensor of the invention may be constructed and arranged in any appropriate manner to carry out the sensing of the target gas species in a gaseous environment containing or susceptible to the presence of same. The fabrication and manufacture of electrochemical cell gas sensors is more fully described in Advances in Electrochemistry and Electrochemical Engineering, Volume 10, John Wiley & Sons, 1976, and the electrochemical cell gas sensor assembly in accordance with the invention may include a chemically selective filter for the electrochemical gas sensing cell, to remove interferent gas components that would otherwise interfere with or preclude measurement of the target gas species in the gas environment being monitored by the gas sensor assembly.

For example, the electrochemical cell gas sensor may be constructed and arranged to electrochemically reduce the gas species to be monitored, so that the monitoring operation involves chemical reduction of the target gas species. Alternatively, the electrochemical cell gas sensor may operate by electrochemically oxidizing the target gas species sought to be detected. As a still further alternative, the electrochemical cell gas sensor may operate by indirect oxidation or reduction reaction of a compound that is produced in the gas sensor device, involving the target gas to be detected in the gas environment being monitored for the presence of the target gas species.

The electrochemical cell gas sensor may be configured to utilize a gas sensor cell containing three electrodes—the working electrode, the reference electrode and the counter electrode, although gas sensor cells having 2-electrode and 4-electrode structures may alternatively be employed. The electrodes are mounted within a housing containing the ionic liquid electrolyte, with the electrodes being coupled in circuit-forming relationship with contacts and electrical wires forming the electronic circuitry of the sensor.

The superiority of the ionic liquid electrolyte medium in the electrochemical gas sensor of the present invention has been demonstrated by relative weight loss tests in which propylene carbonate, a widely used electrolyte of prior art electrochemical gas sensors, and ionic liquids of the present invention (e.g. methyl-octyl-imidazolium-chloride, butyl-methyl-imidazolium-bis-trifluoromethane-sulfonimide) were evaluated at 80° C. After only 5 days at such temperature of 80° C., propylene carbonate, having a boiling point of 240° C., was totally evaporated, while the ionic liquids of the

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present invention were of unchanged mass within the accuracy of the measurement apparatus. These results evidence an unexpected advantage and superiority of the ionic liquids of the present invention over electrolyte characteristic of prior art electrochemical cell gas sensors.

In general, useful electrolytes for electrochemical cells must have an electrical conductivity of at least $10^{-5} \text{ ohm}^{-1} \text{ cm}^{-1}$. The electrical conductivity of the ionic salts of the present invention are typically on the order of $0.01 \text{ ohm}^{-1} \text{ cm}^{-1}$, three orders of magnitude above such electrolytic threshold, providing unexpectedly superior performance of electrochemical cell gas sensors utilizing same. In general, ionic liquids which are preferred for use as electrolyte in the practice of the present invention have an electrical conductivity of at least $10^{-3} \text{ ohm}^{-1} \text{ cm}^{-1}$ and such ionic liquids most preferably include those having an electrical conductivity of at least $10^{-2} \text{ ohm}^{-1} \text{ cm}^{-1}$.

The invention therefore provides an electrochemical cell gas sensor including a gas sensor cell in which an 'ionic liquid' is employed as an electrolyte, e.g., a salt having a melting point below 100° C . and preferably a liquid at room temperature. In contrast to prior art liquids used as electrolyte media, the ionic liquid systems of the invention are characterized by negligible vapour pressure of the electrolyte, and an extremely high electrolytic character since the ionic liquids are constituted by ions.

In another aspect, the invention provides electrochemical fuel cells using ionic liquids as solvent/electrolyte media.

Fuel cells are electrochemical cells consisting of two electrodes connected with an ion-conducting electrolyte. The electrodes usually have a porous structure and are designed to allow very close contact between a gas (the fuel), the electrolyte and the catalyst powder that catalyses the chemical reactions. Oxidation and reduction are separated so that they occur at different electrodes, and the electric circuit is closed by ion transport from one electrode through the electrolyte to the other electrode.

The fuel cells of the present invention in one aspect thereof may be fabricated with cathode and anode elements in an assembly in which the ionic liquid electrolyte medium is disposed between the respective electrodes, in an electrolyte/electrode assembly, with the respective cathode and anode elements have backing layers on their outer surfaces providing pathways for gas access to the respective electrode. On the outer surfaces of the respective backing layers are provided current collector plates including a cathode current collector plate on the cathode-backing layer, and an anode current collector plate on the anode-backing layer. The current collectors are in turn coupled with circuitry including load or output device(s) that utilize the electrical output of the fuel cell.

In contrast to conventional fuel cells that utilize electrolytes that highly restrict the operating range of temperature of the fuel cell (e.g., phosphoric acid fuel cells (PAFCs), using phosphoric acid as electrolyte and operated at about 200° C ; molten carbonate fuel cells (MCFC), using molten carbonates as electrolyte and operated at 500 to 600° C ; solid polymer membrane fuel cells (SPMFC), using conducting polymers as electrolyte and operated below 80° C ; alkaline fuel cells (AFC), using concentrated alkali brine and operated below 100° C ; and solid oxide fuel cells (SOFC) operated at temperatures close to 1000° C), fuel cells utilizing ionic liquid electrolytes in accordance with the present invention can operate at substantially lower temperatures and over significantly wider temperature ranges in which the performance of the prior art fuel cells would be deficient or even non-existent.

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The ionic liquids used as electrolyte media in fuel cells of the present invention may be of any suitable type, including the ionic liquid electrolytes described hereinabove in the preceding discussion of electrochemical cell gas sensors utilizing ionic liquid electrolytes in accordance with the present invention.

The features and advantages of the present invention are more fully appreciated with respect to the following embodiments, described in reference to the drawings.

FIG. 1 is a schematic representation of a process system 10 employing an electrochemical gas sensor according to the present invention, in an illustrative embodiment thereof.

The FIG. 1 process system 10 includes a supply 12 of a source gas. The supply 12 may include a process unit that generates the target gas (i.e., the gas component to be monitored by the electrochemical cell gas sensor) in mixture with other gas components, as a multicomponent gas mixture. Alternatively, the supply 12 of the source gas may be a gas environment that is subject to ingress or contamination by the target gas, e.g., as a toxic, or otherwise hazardous or undesirable gas species in the particular environment. The source gas, containing the target gas as a component thereof, flows from supply 12 in line 14 to the abatement processing unit 16 in which the source gas is treated to remove the target gas therefrom.

A target gas-depleted stream is discharged from the abatement-processing unit 16 in line 18, and may be passed to a further downstream process or final disposition, as required.

A side stream of the source gas from line 14 is flowed in line 20, under the action of motive fluid driver 22, through dust filter 23, interferent species filter 24 and gas sensor 26, being returned to line 14 downstream of gas sensor 26, as shown. The dust filter 23 removes particulates from the source gas, and the interferent species filter 24 removes gas components from the dust-depleted source gas that may interfere with the accurate sensing of the target gas by the electrochemical cell gas sensor. By this arrangement, an interferent-free gas composition, including the target gas species, is flowed from the interferent species filter 24 to the electrochemical cell gas sensor 26.

The electrochemical cell gas sensor 26 monitors the concentration of the target gas species in the side stream and generates a corresponding response signal correlative to the sensed concentration of the target gas species. The response signal is transmitted in signal transmission line 28 to central processing unit (CPU) 30, which in turn generates a corresponding control signal that is transmitted in control signal line 32 to the abatement-processing unit 16. The control signal in line 32 may be employed to modulate the gas processing operation in abatement processing unit 16 to abate the target gas species.

As an illustrative example, if phosgene were the target gas species in the source gas, and such target gas species is abated by chemical reaction thereof with a chemical reagent in the abatement processing unit 16, the amount of the chemical reagent may be modulated in response to the sensed concentration of the phosgene in the source gas, to effect substantially complete removal of the phosgene from the gas stream treated in abatement processing unit 16. In other abatement operations, the process conditions (e.g., temperatures, pressures, flow rates, retention time, etc.) in the abatement processing unit 16 may be modulated to effect the desired reduction in the concentration of the target gas species in the effluent stream being treated.

FIG. 2 is a schematic representation of an electrochemical cell gas sensor 50 according to one embodiment of the invention.

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The electrochemical cell gas sensor **50** comprises a housing **52** formed of a suitable material of construction, e.g., metal, ceramic, polymer, etc. defining therewithin an interior volume. The interior volume of the housing includes an electrolyte compartment **53** containing an ionic liquid electrolyte in accordance with the invention, and an electrode assembly including a counter electrode **54**, a reference electrode **58** and a working electrode **62**. The counter and reference electrodes are separated by separator member **56**, and the reference and working electrodes are separated by separator member **60**.

Overlying the electrode assembly is an interferent species filter **64** for removing interferent species from the source gas flowed therethrough. A dust filter **66** is joined to the housing **52** at the upper end of the housing walls, as shown, being sealed to the top edges of the walls by bond **68**. The bond **68** is formed of a suitable adhesive or sealant medium, and joins the dust filter **66** to the housing **52** in a leak-tight manner, so that source gas flowed through the filter enters the interferent species filter **64** and is prevented from bypassing the filtration and sensing elements in the housing interior volume.

It will be recognized that the electrochemical cell gas sensor **50** is schematically illustrated for ease of description, and does not show the electrical leads to the electrode elements in the housing or other ancillary structure, but based on such description, the electrochemical cell gas sensor **50** may be readily constructed by those skilled in the art, to effect gas sensing operation that is accurate and reproducible for monitoring of the target gas species in the source gas.

FIG. 3 is a simplified schematic representation of an electrochemical fuel cell **100** according to another embodiment of the invention.

The electrochemical fuel cell **100** as illustrated includes an ionic liquid electrolyte compartment **108** containing an ionic liquid electrolyte **102** in accordance with the present invention, and arranged so that the ionic liquid electrolyte in the electrolyte compartment **108** is in contact with anode **106** and cathode **104**.

On the outer surface of the anode (the main surface opposite the surface in contact with the electrolyte) is disposed an anode backing layer **116**. In like manner, the outer surface of the cathode **104** has disposed in contact therewith a cathode-backing layer **110**.

Outwardly of the backing layers **116** and **110** is provided an anode current collector **118** in contact with the anode backing layer **116**, and a cathode current collector **112** in contact with the cathode backing layer **110**, as shown.

The backing layers **116** and **110** are formed of suitable material having pathways therein for gas access to the corresponding electrode with which the backing layer is in contact.

The anode current collector **118** is formed with fuel flow passages **120** therein, accommodating flow of fuel into the passages **120** in the direction indicated by arrow C, with the spent/unused fuel being discharged from such passages in the direction indicated by arrow D. The fuel may comprise hydrogen or any other fuel medium.

The cathode current collector **112** is correspondingly formed with oxidant flow passages **114** therein, into which oxidant is flowed in the direction indicated by arrow A in FIG. 3, and from which spent/unused oxidant is discharged in the direction indicated by arrow B. The oxidant may be of any suitable type, e.g., oxygen, air, oxygen-enriched air, ozone, etc.

The anode and cathode current collectors **118** and **112** in the FIG. 3 fuel cell are coupled with electrical circuitry, schematically represented in FIG. 3 by circuit wire **122**, interconnecting the fuel cell with a load or output device **124** of desired type.

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The fuel cell structure shown in FIG. 3 is highly simplified in character for purposes of illustration, and it will be appreciated that fuel cell apparatus in accordance with the present invention may be fabricated and arranged in a wide variety of conformations and embodiments. The fuel cell apparatus may include a fuel cell stack utilizing a plurality of fuel cell unit structures as shown in FIG. 3, joined together with bipolar current collector plates and end plates, as is known in the art for achieving economies of scale in use of fuel cell elements.

The features, aspects and advantages of the present invention are further shown with reference to the following non-limiting examples relating to the invention.

EXAMPLES

Example 1

An electrochemical cell gas sensor, comprising a gas sensor such as schematically described above, allows for the measurement of hydrogen sulfide (H_2S) concentrations in a gas mixture. The sensor comprises a measuring electrode made of silver (Ag), a reference electrode made of silver (Ag), and a counter electrode made of silver (Ag). Further, the sensor comprises an electrolyte consisting of an ionic liquid, namely ethyl-methyl-imidazolium-trifluoromethane-sulfonate. Concentrations of 20-ppm hydrogen sulfide give rise to a signal current of 0.01 mA. The response time is typically less than 30 s after initial exposure to the gas.

Example 2

An electrochemical cell gas sensor, comprising a gas sensor such as schematically described above, allows for the measurement chlorine (Cl_2) concentrations in a gas mixture. The sensor comprises a measuring electrode made of gold (Au), a reference electrode made of platinum (Pt), and a counter electrode made of platinum (Pt). Further, the sensor comprises an electrolyte consisting of an ionic liquid, namely butyl-methyl-imidazolium-bis-trifluoromethane-sulfonimide. Concentrations of 1-ppm chlorine give rise to a signal current of 0.2 μA . The response time is typically 30 s after initial exposure to the gas.

Although the invention has been variously disclosed herein with reference to illustrative embodiments and features, it will be appreciated that the embodiments and features described hereinabove are not intended to limit the invention, and that other variations, modifications and other embodiments will suggest themselves to those of ordinary skill in the art. The invention therefore is to be broadly construed, consistent with the claims hereafter set forth.

What is claimed is:

1. An electrochemical fuel cell comprising a substantially solvent-free, ionic liquid electrolyte contained within an electrolyte compartment, cathode and anode electrode elements in contact with the ionic liquid electrolyte, and fuel and oxidant passages, wherein the ionic liquid electrolyte is weight-stable after remaining 5 days at a temperature of 80° C. and wherein the substantially solvent-free, ionic liquid electrolyte is substantially free of additives.

2. The electrochemical fuel cell according to claim 1, wherein the ionic liquid electrolyte comprises at least one salt having a melting point less than about 100° C.

3. The electrochemical fuel cell according to claim 1, wherein the ionic liquid electrolyte comprises one salt.

4. The electrochemical fuel cell according to claim 1, wherein the ionic liquid electrolyte comprises a mixture of salts.

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5. The electrochemical fuel cell according to claim 1, wherein the ionic liquid electrolyte has a negligible vapor pressure.

6. The electrochemical fuel cell according to claim 1, wherein the ionic liquid electrolyte comprises at least one salt with the general formula C^+A^- wherein C^+ comprises a cation selected from the group consisting of an imidazolium cation and a pyridinium cation and A^- comprises an anion selected from the group consisting of a halide, a nitrate, a nitrite, a tetrafluoroborate, a hexafluorophosphate, a polyfluoroalkanesulfonate, a bis(trifluoromethylsulfonyl)imide, a methylsulfate, an acetate, and an anion of fluoroalkanoic acid.

7. The electrochemical fuel cell according to claim 1, wherein the ionic liquid electrolyte is selected from the group consisting of methyl-octyl-imidazolium-chloride, and butyl-methyl-imidazolium-bis-trifluoromethane-sulfonimide.

8. The electrochemical fuel cell according to claim 1, wherein the cation includes alkyl and/or aryl substituents, or alkyl and/or aryl substituents further substituted with a substituent selected from the group consisting of a halo, a lower alkyl (C1-C4), a hydroxyl, and an amine.

9. The electrochemical fuel cell according to claim 1, wherein the ionic liquid electrolyte has an electrical conductivity of at least $10^{-3} \text{ ohm}^{-1} \text{ cm}^{-1}$.

10. The electrochemical fuel cell according to claim 1, wherein the ionic liquid electrolyte has an electrical conductivity of at least $10^{-2} \text{ ohm}^{-1} \text{ cm}^{-1}$.

11. The electrochemical fuel cell according to claim 1, wherein the ionic liquid electrolyte further comprises an aluminum halide.

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12. An electrochemical fuel cell, comprising an ionic liquid electrolyte compartment comprising a substantially solvent-free, ionic liquid electrolyte, and an anode and cathode in contact with the ionic liquid electrolyte in the compartment, wherein the anode has an anode backing layer thereon providing passageways therein for gas access to the anode, wherein the cathode has a cathode backing layer thereon providing passageways therein for gas access to the cathode, an anode current collector in contact with the anode backing layer, a cathode current collector in contact with the cathode backing layer, and an electrical circuit-forming means coupled with the anode current collector and the anode current collector, to form therewith an electrical circuit enabling output of electrical energy from the electrochemical fuel cell, wherein the ionic liquid electrolyte is weight-stable after 5 days at a temperature of 80° C .

13. The electrochemical fuel cell of claim 12, wherein the anode current collector has fuel flow passages therein, for flow of fuel therethrough in operation of the electrochemical fuel cell assembly.

14. The electrochemical fuel cell of claim 13, wherein the cathode current collector has oxidant flow passages therein, for flow of oxidant therethrough in operation of the electrochemical fuel cell.

15. The electrochemical fuel cell of claim 12, wherein the electrochemical fuel cell assembly comprises a fuel cell stack assembly.

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